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In the Claims:

1.(currently amended) A process for converting a copper sulfide matte to blister copper, wherein the process includes the steps of:

adding the copper sulphide sulfide matte and flux to a suitable an agitated <u>iron</u> based silicate slag phase; and

injecting, from a discharge tip at the lower end of a top-submerged lance, an oxidizing gas suitable for reacting with the matte to produce blister copper which forms or adds to a continuous blister copper phase below the <u>iron based silicate</u> slag phase;

wherein the lance tip is located within the slag phase at a depth enabling the injected gas to agitate the slag phase, and to react with copper sulphide matte dispersed therein, while precluding a substantial proportion of the gas from contacting the continuous blister copper phase the depth of the iron based silicate slag phase and the level at which the lance tip is located in the slag phase are such that the injected gas agitates the slag phase and reacts with copper sulfide matte dispersed in the slag phase but so that a jet or stream of the injected gas is unable to pass through the lower surface of the slag phase and the gas is substantially precluded from contacting the continuous copper phase.

2.(canceled)

3.(currently amended) The process of claim—2_1, wherein the slag phase has a depth of from about 500 mm up to about 2 m.

4 (currently amended) The process of claim-2_1, wherein the slag phase has a depth of from about 700 mm to about 1.7 m.

5.(previously amended) The process of claim 1, wherein the injection is at a midregion of the height of the slag phase. US Serial No. 10/599570 Page 4 of 15

6.(previously amended) The process of claim 1, wherein the injection is near the upper surface of the slag phase.

7.(currently amended) The process of claim 1, wherein the injection is directed downwardly and laterally outwardly for agitating substantially the entire slag phase for substantially uniform dispersal of the copper sulphide sulfide matte throughout the slag phase.

8.(currently amended) The process of claim 1, wherein the slag phase comprises an-iron based silicate slag is a lime modified iron silicate slag.

9.(original) The process of claim 8, wherein the iron based silicate slag is a ferrous calcium silicate olivine slag.

10.(original) The process of claim 8, wherein the iron based silicate slag is an iron silicate fayalite slag.

11.(currently amended) The process of claim 8, wherein the iron based silicate slag has a ratio of Fe to Si02 of <u>rom-from 1.14</u> to 2.11, a <u>weight to weight ratio of CaO to Fe</u> of from 0.15 to 0.92, and a <u>weight to weight ratio</u> of CaO to Si02 of from 0.22 to 1.11.

12.(currently amended) The process of claim 11, wherein the <u>weight to weight ratio</u> of Fe to Si02 is from 1.14 to 1.55.

13.(original) The process of claim 1, wherein a reductant is added to the slag phase for reducing the formation of magnetite and thereby suppressing foaming of the slag phase.

14.(original) The process of claim 13, wherein the reductant is lump coal.

15.(canceled)

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